

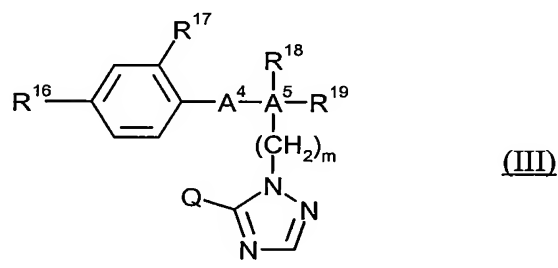
Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application.

Claims 1-3. (Canceled)

4. (Currently amended) ~~Active compound combinations according to Claim 1~~ A composition comprising the carboxamide (1-1) *N*-(3',4'-dichloro-5-fluoro-1,1'-biphenyl-2-yl)-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carboxamide (group 1) and at least one active compound ~~selected from the following groups (2) to (23) according to Claim 1~~

of formula (III)



wherein

Q is hydrogen or SH,

m is 0 or 1,

R¹⁶ is hydrogen, fluorine, chlorine, phenyl or 4-chlorophenoxy,

R¹⁷ is hydrogen or chlorine,

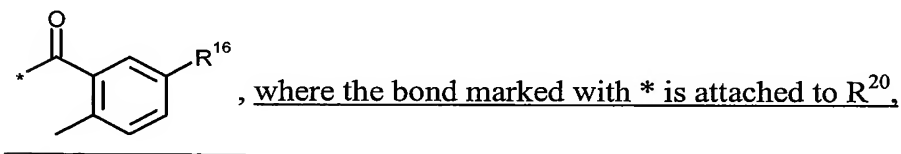
A⁴ is a direct bond, -CH₂-, -(CH₂)₂- or -O-,

R¹⁸ is hydrogen, hydroxyl or cyano, or

A⁴ is *-CH₂-CHR²⁰- or *-CH=CR²⁰- where the bond marked with * is attached to the phenyl ring, in which case

R¹⁸ and R²⁰ together are -CH₂-CH₂-CH[CH(CH₃)₂]- or -CH₂-CH₂-C(CH₃)₂-, or

A⁴ is -N(R²⁰)- and A⁵ together with R¹⁸ and R¹⁹ is the group C=N-R²¹, in which case R²⁰ and R²¹ together are the group



A⁵ is C or Si (silicon),

R¹⁹ is 1-cyclopropylethyl, 1-chlorocyclopropyl, C₁-C₄-alkyl, C₁-C₆-hydroxyalkyl, C₁-C₄-alkylcarbonyl, C₁-C₂-haloalkoxy-C₁-C₂-alkyl, trimethylsilyl-C₁-C₂-alkyl, monofluorophenyl or phenyl, or

R¹⁸ and R¹⁹ together are -O-CH₂-CH(R²¹)-O-, -O-CH₂-CH(R²¹)-CH₂-, or -O-CH(2-chlorophenyl)-, and

R²¹ is hydrogen, C₁-C₄-alkyl or bromine.

Claims 5-12. (Canceled)

13. (Currently amended) ~~Method~~ A method for controlling unwanted phytopathogenic fungi, ~~characterized in that active compound combinations comprising applying the composition according to Claim [[1]] 4 are applied to the unwanted phytopathogenic fungi and/or their habitat.~~
14. (Currently amended) ~~Process~~ A process for preparing fungicidal compositions, ~~characterized in that active compound combinations comprising mixing the composition according to Claim [[1]] 4 are mixed with extenders and/or surfactants.~~
15. (New) The composition according to claim 4, wherein the triazoles of formula III are selected from a group consisting of
- (3-1) azaconazole,
 - (3-2) etaconazole,
 - (3-3) propiconazole,
 - (3-4) difenoconazole,
 - (3-5) bromuconazole,
 - (3-6) cyproconazole,
 - (3-7) hexaconazole,
 - (3-8) penconazole,
 - (3-9) myclobutanil,

- (3-10) tetraconazole,
- (3-11) flutriafol,
- (3-12) epoxiconazole,
- (3-13) flusilazole,
- (3-14) simeconazole,
- (3-15) prothioconazole,
- (3-16) fenbuconazole,
- (3-17) tebuconazole,
- (3-18) ipconazole,
- (3-19) metconazole,
- (3-20) triticonazole,
- (3-21) bitertanol,
- (3-22) triadimenol,
- (3-23) triadimefon,
- (3-24) fluquinconazole, and
- (3-25) quinconazole.

16. (New) The composition according to claim 4, wherein the triazoles of formula III are selected from the group consisting of

(3-3) propiconazole,

(3-12) epoxiconazole,

(3-15) prothioconazole,

(3-17) tebuconazole, and

(3-21) bitertanol.